

Hepatitis C virus (HCV) NS5B polymerase inhibitor

BI-0588

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Summary

BI-0588 is one of the most potent and specific allosteric inhibitors of HCV polymerase known, as demonstrated by its cell-based antiviral replicon activity in the single-digit nM range for genotype(GT)-1. Furthermore, BI-0588 was optimized to combine good *in vitro* metabolic stability, acceptable cell-permeability and good solubility (@ pH 7). Following *in vivo* cross-species PK profiling, this compound was further progressed into preclinical drug development. Hence, the molecule is suitable for *in vitro* as well as *in vivo* experiments.

Chemical Structure

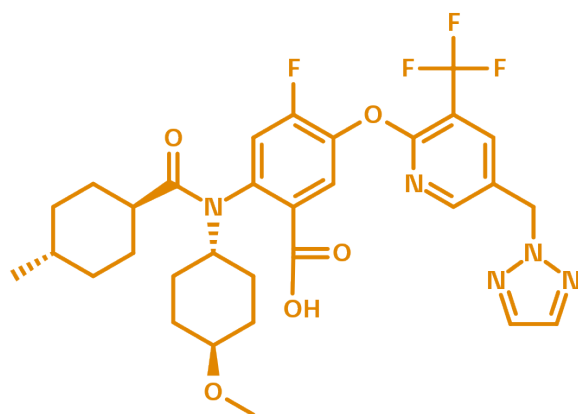


Figure 1: 2D structure of BI-0588

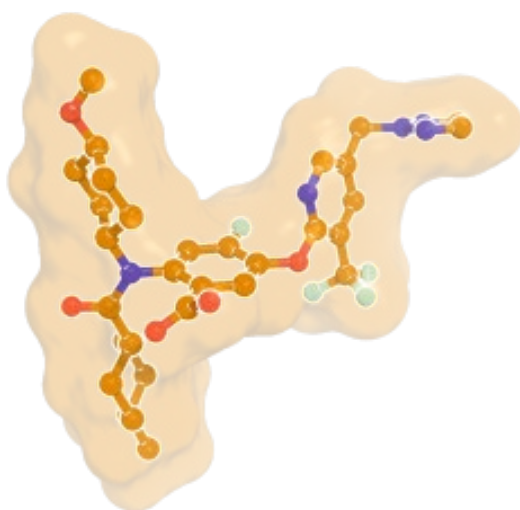


Figure 2: BI-0588, 3D conformation

Highlights

BI-0588 is a highly potent and selective allosteric thumb pocket-2 inhibitor of Hepatitis C virus (HCV) NS5B polymerase and complements our allosteric thumb pocket-1 inhibitor, [deleobuvir](#). It has demonstrated optimized antiviral activity in a cell-based replicon system, with activity in the low nanomolar range for genotype-1 HCV replication. Its overall CMC-ADME-PK profile is good. BI-0588 is also suitable for both *in vitro* and *in vivo* experiments, as it has shown sufficient plasma exposure in rodents and was found to predominantly localize to the liver.

Target information

HCV NS5B is an RNA-dependent RNA polymerase that is essential for the replication of the genome of the hepatitis C virus. BI-0588 inhibits the polymerase activity by binding to the allosteric pocket-2 located in the thumb domain of NS5B⁵.

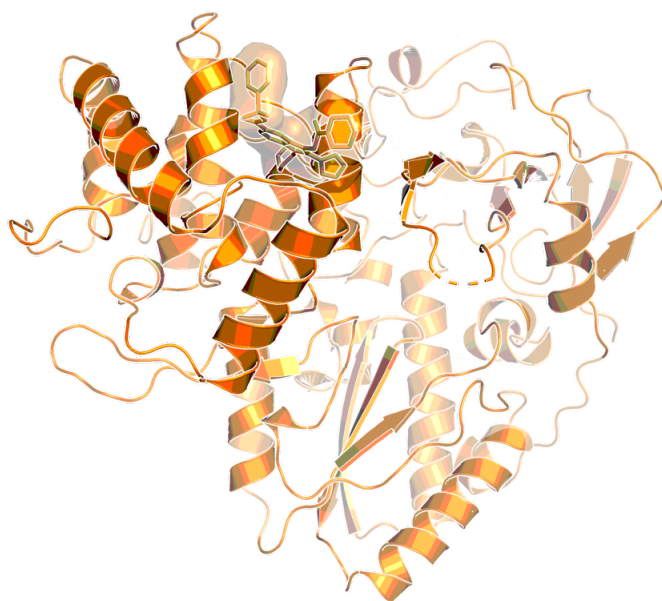


Figure 3: Binding model of BI-0588 on basis of an X-ray structure with a related inhibitor (PDB code: 4JJS)

In vitro activity

BI-0588 is a potent and specific allosteric pocket-2 inhibitor of the virally-encoded NS5B polymerase, and inhibited genotype (GT)-1 HCV replication in the cell-based replicon system in the single-digit nM range with $EC_{50} = 2.3$ nM and $EC_{50} = 3.0$ nM for GT1a and GT1b, respectively.

PROBE NAME / NEGATIVE CONTROL	BI-0588	BI-0900
MW [Da, free base] ^a	633.6	551.5
HCV-POLd21 (IC ₅₀) [nM] ^b	60	6,400
HCVLUC1 (EC ₅₀) [nM] ^c	3.3	>10,000
Rep GT 1a/1b (EC ₅₀) [nM] ^d	2.3 / 3.0	n.d.

^a For the salt form you will get, please refer to the label on the vial and for the molecular weight of the salt, please refer to the FAQs

^b Scintillation proximity HCV polymerase assay with recombinant protein His-NS5Bd21. This assay measures the incorporation of 3H-UTP during the elongation of RNA primer

^c Cell-based luciferase reporter HCV RNA replication assay, MP-1 cells with 10% FBS, 28 h incubation

^d Cell-based HCV replicon assay using RT-PCR for RNA quantification, genotype background 1a and 1b, Huh7 cells, 72 h incubation

In vitro DMPK and CMC parameters

BI-0588 is highly soluble at neutral pH while being cell-permeable. A hallmark is its good metabolic stability in hepatocyte across several species.

PROBE NAME / NEGATIVE CONTROL	BI-0588	BI-0900
logD @ pH 7.4	1.8	0.8
Solubility @ pH 7 [µg/mL]	193	ongoing
Caco-2 permeability AB @ pH 7.4 [$\times 10^{-6}$ cm/s]	3.0	0.1
Caco-2 efflux ratio	2.7	102.8
Microsomal stability (human) [% Q _H]	<24	<23
Hepatocyte stability (human / dog / rat) [% Q _H]	<12 / <19 / 10	n.a.
Plasma Protein Binding (human / dog / rat) [%]	99.6 / 99.5 / 99.8	n.a.
hERG [inh. % @ 10 µM]	1.9%	n.a.
CYP 3A4 (IC ₅₀) [µM]	8.1	>50
CYP 2C9 (IC ₅₀) [µM]	6.5	>50
CYP 2C19 (IC ₅₀) [µM]	9.4	>50

CYP 2D6 (IC ₅₀) [μM]	>30	n.a.
CYP 1A2 (IC ₅₀) [μM]	>30	n.a.

In vivo DMPK parameters

BI-0588 demonstrates sufficient plasma exposure in rat and dog species. As often observed for this anthranilic acid chemical series, BI-0588 illustrates that *in vitro* hepatic clearance was not predictive of the *in vivo* clearance, and biliary excretion of the parent compound significantly contributes to clearance mechanism.

BI-0588	DOG	RAT
Clearance [mL/(min*kg)] ^a	56	44
Mean residence time after <i>i.v.</i> dose [L/kg] ^a	1.0	3.8
t _{1/2} [h]	2.1	6.2
F [%]	44	41
V _{ss} [L/kg]	1.2	6.4

^a *i.v.* dose: 1.2mg/kg, *p.o.* dose: 3.1 mg/kg

In vivo pharmacology

During the course of this drug development project, the lack of a small animal model to study HCV biology and virus-host interaction imposed an additional challenge.

Boehringer Ingelheim's scientists developed a novel *in vitro-in vivo* proportionality method to assess a human **Liver Corrected Inhibitory Quotient** (LCIQ)³ predictive of clinical efficacy. Given that Hepatitis-C virus replicates predominantly in the human liver, the main objective was to predict the local drug concentration – i.e. drug's concentration within the human liver at C_{min} – and the required dose to achieve such local exposure.

Plasma exposure and liver concentration were measured in rodents, and then coupled with allometry scaling to guide candidate selection. From our studies the research target profile was determined: LCIQ >500 [(plasma C_{min}) / EC₅₀]*Kp_(liver) >500]. Based on the PK profile.

The Human Dose Prediction to achieve LCIQ500 was: 357 mg (bid)

[Rep Gt1b EC₅₀: 0.003 μM; liver Kp: 78; estimated C_{min}: 0.02 μM; predicted C_{max}: 0.80 μM]

We found that BI-0588 localizes predominantly into the liver where HCV mainly replicates, given its high liver partitioning coefficient ([liver]/[plasma]= Kp_{liver}: 78 in rodents).

Negative control

Despite close structural similarity, BI-0900 bears a tetrahydropyran (THP) ring instead of a 4-methylcyclohexane ring, thus imparting beneficial interaction in the deep lipophilic subpocket. This translates into a 107-fold and >3000-fold potency shift in HCV-POLd21 and HCV Luc1 assays, respectively.

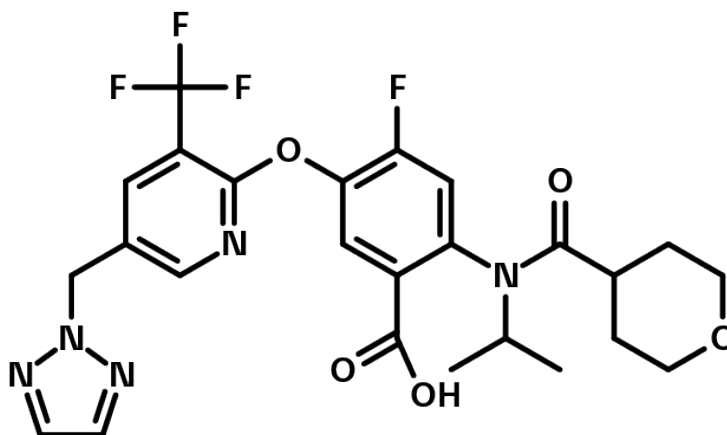



Figure 4: BI-0900 which serves as a negative control

Selectivity

SELECTIVITY DATA AVAILABLE	BI-0588	BI-0900
SafetyScreen44™ with kind support of  eurofins	Yes	Yes
Invitrogen®	No	No
DiscoverX®	No	No
Dundee	No	No

Co-crystal structure of the Boehringer Ingelheim probe compound and the target protein.

The X-ray crystal structure of the target in complex with a related inhibitor is available (PDB code: 4JJS)².

Reference molecule(s)

See reference 6.

Supplementary data

2D structure files can be downloaded free of charge from [openMe](#).

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